Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	4353	((514/679) or (514/721) or (514/880) or (514/881) or (514/901) or (424/49) or (424/76.8)).CCLS.	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	OFF	2005/11/22 10:26
L2	1507	hydroxydiphenyl near2 ether	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/11/22 10:26
L3	171	l1 and l2	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/11/22 10:01
L4	81762	antimicrob$4	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/11/22 10:26
L5	128	l3 and l4	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/11/22 10:01
L6	6865	antimicrob$4.clm.	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/11/22 10:26
L7	143	hydroxydiphenyl near2 ether.clm.	US-PGPUB; USPAT; USOCR	OR	ON	2005/11/22 10:26
L8	4104	((514/679) or (514/721) or (514/880) or (514/881) or (514/901) or (424/49) or (424/76.8)).CCLS.	US-PGPUB; USPAT; USOCR	OR	OFF	2005/11/22 10:26
L9	10	l6 and l7 and l8	US-PGPUB; USPAT; USOCR; EPO; JPO; DERWENT	OR	ON	2005/11/22 10:27

10/816,967

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PASSWORD:
TERMINAL (ENTER 1, 2, 3, OR ?):2

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NEWS	1		Web Page URLs for STN Seminar Schedule - N. America
NEWS	2		"Ask CAS" for self-help around the clock
NEWS	3	SEP 09	ACD predicted properties enhanced in REGISTRY/ZREGISTRY
NEWS	4	OCT 03	MATHDI removed from STN
NEWS	5	OCT 04	CA/CAplus-Canadian Intellectual Property Office (CIPO) added to core patent offices
NEWS	6	OCT 06	STN AnaVist workshops to be held in North America
NEWS	7	OCT 13	New CAS Information Use Policies Effective October 17, 2005
NEWS	8	OCT 17	STN(R) AnaVist(TM), Version 1.01, allows the export/download of CAplus documents for use in third-party analysis and visualization tools
NEWS	9	OCT 27	Free KWIC format extended in full-text databases
NEWS	10	OCT 27	DIOGENES content streamlined
NEWS	11	OCT 27	EPFULL enhanced with additional content
NEWS	12	NOV 14	CA/CAplus - Expanded coverage of German academic research

NEWS EXPRESS NOVEMBER 18 CURRENT VERSION FOR WINDOWS IS V8.01,
CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
AND CURRENT DISCOVER FILE IS DATED 13 JUNE 2005.
V8.0 USERS CAN OBTAIN THE UPGRADE TO V8.01 AT
http://download.cas.org/express/v8.0-Discover/

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* * * * * * * * * * * * * STN Columbus * * * * * * * * * * * * * * * *

FILE 'HOME' ENTERED AT 08:19:46 ON 22 NOV 2005

=> file reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

10/816,967

FILE 'REGISTRY' ENTERED AT 08:19:57 ON 22 NOV 2005
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STRUCTURE FILE UPDATES: 21 NOV 2005 HIGHEST RN 868586-21-4
DICTIONARY FILE UPDATES: 21 NOV 2005 HIGHEST RN 868586-21-4

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TSCA INFORMATION NOW CURRENT THROUGH JULY 14, 2005

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**********************************************************************
*                                                                    *
* The CA roles and document type information have been removed from *
* the IDE default display format and the ED field has been added,    *
* effective March 20, 2005.  A new display format, IDERL, is now     *
* available and contains the CA role and document type information.  *
*                                                                    *
**********************************************************************
```

Structure search iteration limits have been increased. See HELP SLIMITS
for details.

REGISTRY includes numerically searchable data for experimental and
predicted properties as well as tags indicating availability of
experimental property data in the original document. For information
on property searching in REGISTRY, refer to:

http://www.cas.org/ONLINE/UG/regprops.html

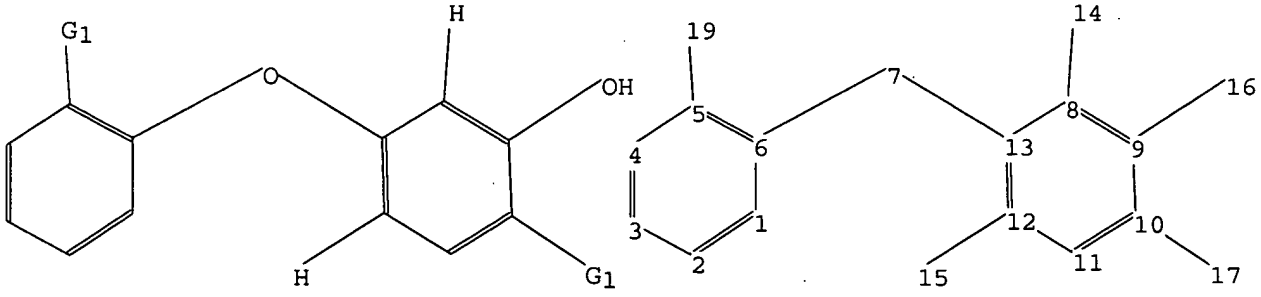
=> ....Testing the current file.... screen

ENTER SCREEN EXPRESSION OR (END):end

=> screen 1992 OR 2016 OR 2021 OR 2026 OR 1929 OR 1840

L1 SCREEN CREATED

=>
Uploading C:\Program Files\Stnexp\Queries\10816967.str



chain nodes :
7 14 15 16 17 19

ring nodes :
1 2 3 4 5 6 8 9 10 11 12 13
chain bonds :
5-19 6-7 7-13 8-14 9-16 10-17 12-15
ring bonds :
1-2 1-6 2-3 3-4 4-5 5-6 8-13 8-9 9-10 10-11 11-12 12-13
exact/norm bonds :
5-19 6-7 7-13 9-16 10-17
exact bonds :
8-14 12-15
normalized bonds :
1-2 1-6 2-3 3-4 4-5 5-6 8-13 8-9 9-10 10-11 11-12 12-13
isolated ring systems :
containing 1 : 8 :

G1:H,n-Pr,i-Pr,n-Bu,i-Bu,s-Bu,t-Bu

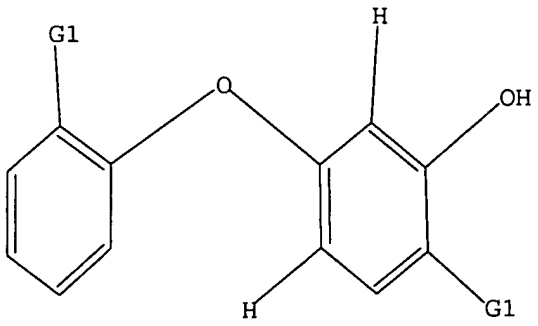
Match level :
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:Atom 7:CLASS 8:Atom 9:Atom 10:Atom
11:Atom 12:Atom 13:Atom 14:CLASS 15:CLASS 16:CLASS 17:CLASS 19:CLASS

L2 STRUCTURE UPLOADED

=> que L2 NOT L1

L3 QUE L2 NOT L1

=> d
L3 HAS NO ANSWERS
L1 SCR 1992 OR 2016 OR 2021 OR 2026 OR 1929 OR 1840
L2 STR



G1 H,n-Pr,i-Pr,n-Bu,i-Bu,s-Bu,t-Bu

Structure attributes must be viewed using STN Express query preparation.
L3 QUE L2 NOT L1

=> s l3
SAMPLE SEARCH INITIATED 08:20:34 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 660 TO ITERATE

100.0% PROCESSED 660 ITERATIONS 2 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 11659 TO 14741
PROJECTED ANSWERS: 2 TO 124

L4 2 SEA SSS SAM L2 NOT L1

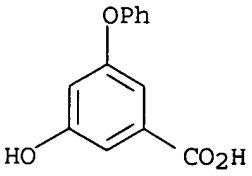
=> s l3 ful
FULL SEARCH INITIATED 08:20:43 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 12922 TO ITERATE

100.0% PROCESSED 12922 ITERATIONS 73 ANSWERS
SEARCH TIME: 00.00.01

L5 73 SEA SSS FUL L2 NOT L1

=> d scan

L5 73 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Benzoic acid, 3-hydroxy-5-phenoxy- (9CI)
MF C13 H10 O4



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):0

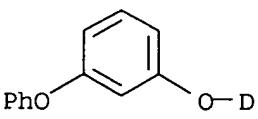
=> s l3 ful css
FULL SEARCH INITIATED 08:21:29 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 12922 TO ITERATE

100.0% PROCESSED 12922 ITERATIONS 10 ANSWERS
SEARCH TIME: 00.00.01

L6 10 SEA CSS FUL L2 NOT L1

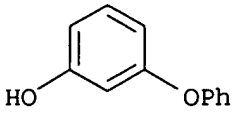
=> d scan

L6 10 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Phenol-d, 3-phenoxy- (9CI)
MF C12 H9 D O2

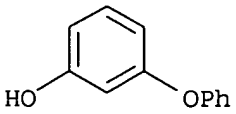


HOW MANY MORE ANSWERS DO YOU WISH TO SCAN? (1):9

L6 10 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Phenol, 3-phenoxy-, potassium salt (9CI)
MF C12 H10 O2 . K

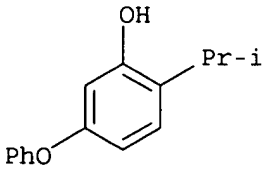


L6 10 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Phenol, 3-phenoxy- (9CI)
MF C12 H10 O2
CI COM



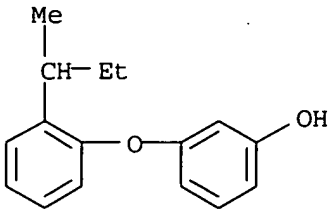
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L6 10 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Phenol, 2-(1-methylethyl)-5-phenoxy- (9CI)
MF C15 H16 O2



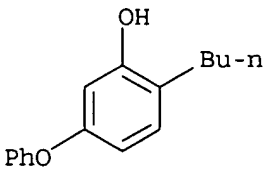
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L6 10 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Phenol, 3-[2-(1-methylpropyl)phenoxy]- (9CI)
MF C16 H18 O2



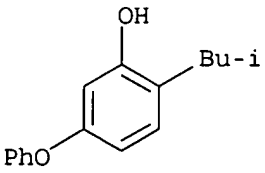
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L6 10 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Phenol, 2-butyl-5-phenoxy- (9CI)
MF C16 H18 O2



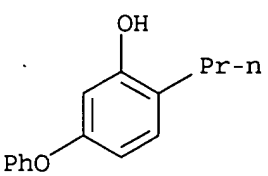
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L6 10 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Phenol, 2-(2-methylpropyl)-5-phenoxy- (9CI)
MF C16 H18 O2



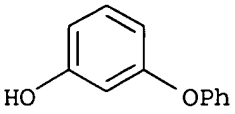
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L6 10 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Phenol, 5-phenoxy-2-propyl- (9CI)
MF C15 H16 O2



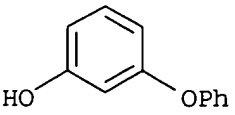
PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

L6 10 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Phenol, 3-phenoxy-, sodium salt (9CI)
MF C12 H10 O2 . Na



● Na

L6 10 ANSWERS REGISTRY COPYRIGHT 2005 ACS on STN
IN Phenol, 3-phenoxy-, barium salt (9CI)
MF C12 H10 O2 . 1/2 Ba



●1/2 Ba

ALL ANSWERS HAVE BEEN SCANNED

=> s l6 and (c15h16o2 or c16h18o2)
29751 C15H16O2
3292 C16H18O2
L7 5 L6 AND (C15H16O2 OR C16H18O2)

=> file caplus
COST IN U.S. DOLLARS SINCE FILE ENTRY TOTAL SESSION
FULL ESTIMATED COST 334.87 335.08

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FILE COVERS 1907 - 22 Nov 2005 VOL 143 ISS 22
FILE LAST UPDATED: 21 Nov 2005 (20051121/ED)

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=> s l7
L8 10 L7

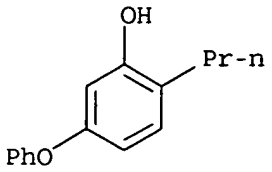
=> dup rem l8
PROCESSING COMPLETED FOR L8
L9 10 DUP REM L8 (0 DUPLICATES REMOVED)

=> d 1-10 bib ab fhitstr

L9 ANSWER 1 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:22652 CAPLUS
DN 138:78170
TI Cosmetic composition comprising a hydroxydiphenyl ether derivative for inhibiting the development of body odors
IN Forestier, Serge; Courbiere, Christophe
PA L'Oreal, Fr.
SO PCT Int. Appl., 22 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003002081	A1	20030109	WO 2002-FR1790	20020528
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	FR 2826574	A1	20030103	FR 2001-8662	20010629
	FR 2826574	B1	20050826		
PRAI	FR 2001-8662	A	20010629		

OS MARPAT 138:78170
AB The invention relates to a cosmetic or dermopharmaceutical composition comprising at least one hydroxydiphenyl ether derivative and furthermore at least one compound selected from active deodorants or antiperspirants. The invention also relates to a method for the treatment of body odors, in particular of the armpit, using the above compns. Formulation of a deodorant stick containing 4,4'-dihydroxydiphenyl ether is disclosed.
IT **194793-00-5**
RL: COS (Cosmetic use); BIOL (Biological study); USES (Uses)
(cosmetic compns. containing hydroxydiphenyl ether derivative for inhibiting body odors)
RN 194793-00-5 CAPLUS
CN Phenol, 5-phenoxy-2-propyl- (9CI) (CA INDEX NAME)

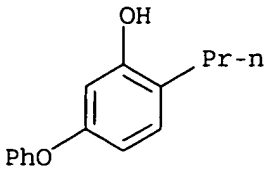


RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 2 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:22651 CAPLUS
DN 138:78169
TI Cosmetic compositions containing a derivative of hydroxydiphenyl ether for inhibiting the development of body odors
IN Forestier, Serge; Courbiere, Christophe
PA L'Oreal, Fr.
SO PCT Int. Appl., 56 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003002080	A1	20030109	WO 2002-FR1789	20020528
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	FR 2826573	A1	20030103	FR 2001-8661	20010629
	FR 2826573	B1	20051007		
PRAI	FR 2001-8661	A	20010629		

OS MARPAT 138:78169
AB The invention relates to a cosmetic composition or dermopharmaceutical composition
comprising at least one hydroxydiphenyl ether derivative and at least one specific conditioning agent. The invention also relates to a method for treating human body odors, particularly axillary odors, using said compns. Formulations of deodorants containing 4,4'-dihydroxydiphenyl ether are disclosed.
IT **194793-00-5**
RL: COS (Cosmetic use); BIOL (Biological study); USES (Uses)
(cosmetic compns. containing derivative of hydroxydiphenyl ether for inhibiting
development of body odors)
RN 194793-00-5 CAPLUS
CN Phenol, 5-phenoxy-2-propyl- (9CI) (CA INDEX NAME)

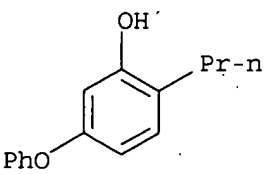


RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 3 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:22650 CAPLUS
DN 138:78168
TI Cosmetic compositions containing a hydroxydiphenyl ether derivative for inhibiting body odors
IN Forestier, Serge; Courbiere, Christophe
PA L'Oreal, Fr.
SO PCT Int. Appl., 43 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2003002079	A1	20030109	WO 2002-FR1787	20020528
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
FR 2826570	A1	20030103	FR 2001-8658	20010629
FR 2826570	B1	20050826		
PRAI FR 2001-8658	A	20010629		

OS MARPAT 138:78168
AB The invention concerns a cosmetic or dermopharmaceutical composition comprising at least a hydroxydiphenyl ether derivative and furthermore at least a particular thickening polymer. The invention also concerns a method for treating human body odors and in particular axillary odors with such compns. Formulations of deodorants containing 4,4'-dihydroxydiphenyl ether are disclosed.
IT **194793-00-5**
RL: COS (Cosmetic use); BIOL (Biological study); USES (Uses)
(cosmetic compns. containing hydroxydiphenyl ether derivative for inhibiting body odors)
RN 194793-00-5 CAPLUS
CN Phenol, 5-phenoxy-2-propyl- (9CI) (CA INDEX NAME)



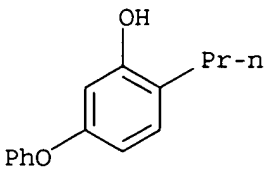
RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 4 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:22649 CAPLUS
DN 138:78167
TI Cosmetic compositions containing a hydroxydiphenyl ether derivative for inhibiting body odors
IN Forestier, Serge; Courbiere, Christophe
PA L'Oreal, Fr.
SO PCT Int. Appl., 22 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2003002078	A1	20030109	WO 2002-FR1786	20020528
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
FR 2826571	A1	20030103	FR 2001-8659	20010629
FR 2826571	B1	20051007		
PRAI FR 2001-8659	A	20010629		

OS MARPAT 138:78167
AB The invention relates to an anhydrous cosmetic or dermopharmaceutical composition or comprising in a lipophilic phase at least a hydroxydiphenyl ether derivative The invention also relates to a method for treating human body odors, particularly axillary odors, using said compns. Formulation of a deodorant containing 4,4'-dihydroxydiphenyl ether is disclosed.
IT **194793-00-5**
RL: COS (Cosmetic use); BIOL (Biological study); USES (Uses)
(cosmetic compns. containing hydroxydiphenyl ether derivative for inhibiting body odors)
RN 194793-00-5 CAPLUS
CN Phenol, 5-phenoxy-2-propyl- (9CI) (CA INDEX NAME)



RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

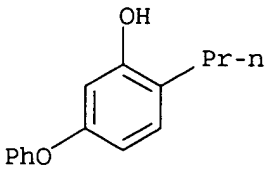
L9 ANSWER 5 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:22643 CAPLUS
DN 138:78166
TI Cosmetic compositions containing a hydroxydiphenyl ether derivative for inhibiting body odors
IN Forestier, Serge; Courbiere, Christophe

10/816,967

PA L'Oreal, Fr.
SO PCT Int. Appl., 23 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2003002072	A1	20030109	WO 2002-FR1788	20020528
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
FR 2826572	A1	20030103	FR 2001-8660	20010629
FR 2826572	B1	20051007		
PRAI FR 2001-8660	A	20010629		

OS MARPAT 138:78166
AB The invention concerns an aerosol device consisting of a container comprising a aerosol composition consisting of a liquid phase (a) (or liquor) comprising at least a hydroxydiphenyl ether derivative and (b) at least a particular propellant and of means for dispensing said aerosol composition as well as the method for treating human body odors and in particular axillary odors with said device. Formulation of a deodorant aerosol containing 4,4'-dihydroxydiphenyl ether 2.0, and ethanol q.s. 100.0 is disclosed.
IT **194793-00-5**
RL: COS (Cosmetic use); BIOL (Biological study); USES (Uses)
(cosmetic compns. containing hydroxydiphenyl ether derivative for inhibiting body odors)
RN 194793-00-5 CAPLUS
CN Phenol, 5-phenoxy-2-propyl- (9CI) (CA INDEX NAME)

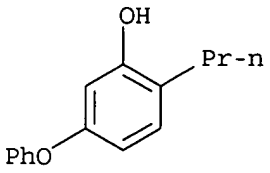


RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 6 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2003:811645 CAPLUS
DN 139:311958
TI Deodorants and antiperspirants especially for men containing hydroxydiphenyl ethers as arylsulfatase inhibitors
IN Banowski, Bernhard; Wadle, Armin; Siegert, Petra
PA Henkel Kgaa, Germany
SO Ger. Offen., 20 pp.
CODEN: GWXXBX
DT Patent
LA German
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

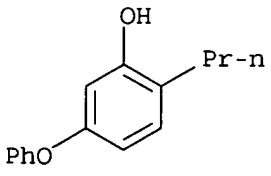
PI DE 10216368 A1 20031016 DE 2002-10216368 20020412
WO 2003086338 A1 20031023 WO 2003-EP3603 20030407
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
EP 1494640 A1 20050112 EP 2003-720431 20030407
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
JP 2005530724 T2 20051013 JP 2003-583362 20030407
US 2005203179 A1 20050915 US 2005-511015 20050422
PRAI DE 2002-10216368 A 20020412
WO 2003-EP3603 W 20030407
OS MARPAT 139:311958
AB The invention concerns deodorant and antiperspirant compns. that contain hydroxydiphenyl ethers as arylsulfatase inhibitors. Arylsulfate inhibition results in the decrease of body odor caused by the decomposition of steroid esters, especially in men; therefore the inhibitors are applied especially in men's deodorants. A water-free, surfactant-containing formulation included (weight/weight%): silicone oil DC 245 28; Eutanol G 16 10; Ucon Fluid AP 5; Cutina HR 6; Lorol C18 20; Eumulgin B3 3; aluminum chlorohydrate 7.995; 4-(2,5-dimethylphenoxy)-phenol 0.005.
IT **194793-00-5**
RL: COS (Cosmetic use); BIOL (Biological study); USES (Uses)
(deodorants and antiperspirants especially for men containing hydroxydiphenyl ethers as arylsulfatase inhibitors)
RN 194793-00-5 CAPLUS
CN Phenol, 5-phenoxy-2-propyl- (9CI) (CA INDEX NAME)



L9 ANSWER 7 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2000:911075 CAPLUS
DN 134:71589
TI Preparation of 5-(halo or alkyl)-5-aryl-2,4-thiazolidinedione and oxazolidinedione derivatives as PPAR agonists
IN Sahoo, Souyma P.; Santini, Conrad; Boueres, Julia K.; Heck, James V.; Metzger, Edward; Lombardo, Victoria K.
PA Merck & Co., Inc., USA
SO PCT Int. Appl., 140 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

PI WO 2000078312 A1 20001228 WO 2000-US16586 20000616
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
CA 2376919 AA 20001228 CA 2000-2376919 20000616
EP 1194146 A1 20020410 EP 2000-944694 20000616
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO
US 6399640 B1 20020604 US 2000-595802 20000616
JP 2003502369 T2 20030121 JP 2001-504375 20000616
AU 773505 B2 20040527 AU 2000-58755 20000616
PRAI US 1999-139953P P 19990618
WO 2000-US16586 W 20000616
OS MARPAT 134:71589
AB The title compds. (I) [wherein Ar1 = (hetero)arylene optionally substituted with 1-4 R1 groups; Ar2 = (hetero)aryl substituted with 1-5 Ra groups; X and Y = independently O, S, NRb, or CH2; Z = O or S; n = 0-3; R = (un)substituted alkyl, F, or Cl; Ra = halo, ORb, (hetero)aryl, or (un)substituted alkanoyl, alkyl, alkenyl, alkynyl, or heterocyclyl; Rb = H, (hetero)aryl, (hetero)arylalkyl, alkanoyl, cycloalkyl, or (un)substituted alkyl, alkenyl, or alkynyl] were prepared as peroxisome proliferator activated receptor (PPAR) agonists. For example, 4-(3-bromopropoxy)-3-propylphenyl Ph ether and Me 3-hydroxyphenylacetate were coupled. The acetate was α-brominated with N-bromosuccinimide and then treated with thiourea and NaOAc in MeOEt to give the 5-aryl-2,4-thiazolidinedione cycloaddn. product. Fluorination with N-fluorobenzenesulfonimide in the presence of KOBu-t in DMF, followed by addition of NaN(TMS)2, afforded the 5-aryl-5-fluoro-2,4-thiazolidinedione (II). I are useful in the treatment, control, or prevention of diabetes, hyperglycemia, hyperlipidemia (including hypercholesterolemia and hypertriglyceridemia), atherosclerosis, obesity, vascular restenosis, and other PPAR α and/or γ mediated diseases, disorders, and conditions (no data).
IT **194793-00-5P**, 2-Propyl-5-phenoxyphenol
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of 5-(halo or alkyl)-5-aryl-2,4-thiazolidinedione and oxazolidinedione PPAR agonists by cycloaddn. of (thio)urea with α-halophenylacetates followed by halogenation or alkylation)
RN 194793-00-5 CAPLUS
CN Phenol, 5-phenoxy-2-propyl- (9CI) (CA INDEX NAME)



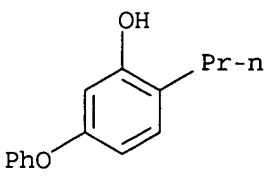
RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 8 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
AN 2000:822696 CAPLUS

DN 133:362617
TI Preparation of hydroxydiphenyl ethers as antimicrobials.
IN Holzl, Werner; Haap, Wolfgang; Ochs, Dietmar; Puchtler, Karin; Schnyder, Marcel; Kulkarni, Surendra Umesh; Radhakrishna, Arakali Srinivasarao; Sawant, Mangesh Shivram; Mahtre, Asawari Bhikaji
PA Ciba Specialty Chemicals Holding Inc., Switz.
SO Eur. Pat. Appl., 42 pp.
CODEN: EPXXDW
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	EP 1053989	A2	20001122	EP 2000-810404	20000511
	EP 1053989	A3	20040121		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	JP 2001011005	A2	20010116	JP 2000-142621	20000516
	CN 1275376	A	20001206	CN 2000-108932	20000519
	BR 2000002441	A	20010102	BR 2000-2441	20000519
	US 2003162836	A1	20030828	US 2002-281011	20021025
	US 2004186174	A1	20040923	US 2004-816967	20040402
PRAI	EP 1999-810442	A	19990520		
	US 2000-573403	A1	20000518		
	US 2002-281011	B1	20021025		

OS MARPAT 133:362617
AB Use of title compds. [I; when the OH is in the para position with respect to the ether linkage, then R1, R2 = H, OH, alkyl, cycloalkyl, alkylcarbonyl, alkoxy, Ph, phenylalkyl; R3 = H, alkyl, alkoxy; R4 = H, alkyl, hydroxyalkyl, cycloalkyl, OH, CHO, acetonyl, alkylcarbonyl, alkenyl, CO2H, carboxyalkyl, alkylcarbonylalkyl, carboxyallyl; when the OH is in the meta position, then R2 = H, alkyl, hydroxyalkyl, alkylcarbonyl; R1, R3 = H, alkylcarbonyl, alkyl; R4 = H, alkyl, hydroxyalkyl, cycloalkyl, OH, CHO, acetonyl, alkylcarbonyl, alkenyl, CO2H, carboxyalkyl, alkylcarbonylalkyl, carboxyallyl; when the OH is ortho, then R1 = H, alkylcarbonyl, alkyl; R4 = H, alkyl, hydroxyalkyl, cycloalkyl, OH, CHO, acetonyl, alkylcarbonyl, alkenyl, CO2H, carboxyalkyl, alkylcarbonylalkyl, carboxyallyl; R2, R3 = H, alkylcarbonyl, alkyl; with provisos] as antimicrobials is claimed. Thus, 2,5-dimethylphenol, 4-bromoanisole, KOH and Cu powder were heated at 160° for 5 h to give 40% 4-(2,5-dimethylphenoxy)anisole. The latter was refluxed 4 h with aqueous HBr in HOAc to give 52% 4-(2,5-dimethylphenoxy)phenol. Tested I showed min. inhibitory concns. of 12.5-25 ppm against Candida albicans ATCC 10231.
IT **194793-00-5P**
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BUU (Biological use, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of hydroxydiphenyl ethers as antimicrobials)
RN 194793-00-5 CAPLUS
CN Phenol, 5-phenoxy-2-propyl- (9CI) (CA INDEX NAME)

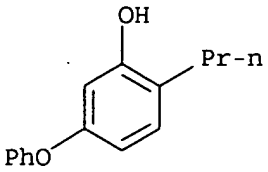


L9 ANSWER 9 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
AN 1999:421669 CAPLUS
DN 131:73645
TI Preparation of arylthiazolidinediones as agonists of peroxisome proliferator activated receptor.
IN Sahoo, Soumya P.; Tolman, Richard L.; Han, Wei; Bergmann, Jeffrey; Santini, Conrad; Lombardo, Vicki R.; Desai, Ranjit; Boueres, Julia K.; Gratale, Dominick F.
PA Merck & Co., Inc., USA
SO PCT Int. Appl., 133 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9932465	A1	19990701	WO 1998-US27139	19981218
	W: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GD, GE, HR, HU, ID, IL, IN, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, SL, TJ, TM, TR, TT, UA, US, UZ, VN, YU, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	US 6008237	A	19991228	US 1998-213542	19981217
	CA 2315397	AA	19990701	CA 1998-2315397	19981218
	AU 9918334	A1	19990712	AU 1999-18334	19981218
	AU 740733	B2	20011115		
	BR 9813801	A	20001003	BR 1998-13801	19981218
	EP 1040102	A1	20001004	EP 1998-963283	19981218
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, SI, LT, LV, FI, RO				
	TR 200001753	T2	20001121	TR 2000-200001753	19981218
	JP 2001526278	T2	20011218	JP 2000-525402	19981218
	JP 3373198	B2	20030204		
	ZA 9903232	A	19991111	ZA 1999-3232	19990511
	NO 2000003112	A	20000818	NO 2000-3112	20000616
	BG 104602	A	20010131	BG 2000-104602	20000713
PRAI	US 1997-68271P	P	19971219		
	GB 1998-16279	A	19980727		
	US 1998-105238P	P	19981022		
	WO 1998-US27139	W	19981218		

OS MARPAT 131:73645
AB Title compds. [I; Ar1 = (substituted) arylene, heteroarylene; Ar2 = o-substituted aryl, heteroaryl; X, Y = O, S, imino, CH2; Z = O, S; n = 0-3], were prepared for treatment of of diabetes, hyperglycemia, hyperlipidemia, atherosclerosis, obesity, vascular restenosis, etc. (no data). Thus, Me 4-hydroxyphenylacetate, Br(CH2)3Br, and K2CO3 were stirred overnight in DMF to give Me 4-(3-bromophenoxy)phenylacetate. This was stirred with 4-phenoxy-2-propylphenol and Cs2CO3 in DMF at 40° overnight to give Me 4-[3-(2-propyl-4-phenoxyphenoxy)propoxy]phenylacetate. The latter was added to a mixture of LiN(SiMe3)2 and Me3SiCl in THF at -78°; after 2 h N-bromosuccinimide was added and the mixture was stirred overnight at room temperature to give the α-bromo derivative, which was stirred with thiourea and NaOAc in methoxyethanol at 115° for 5 h to give 5-[4-[3-(2-propyl-4-phenoxyphenoxy)propoxy]phenyl]-2,4-thiazolidinedione.
IT **194793-00-5P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn of arylthiazolidinedione derivs. as peroxisome proliferator activated receptor agonists)

RN 194793-00-5 CAPLUS
CN Phenol, 5-phenoxy-2-propyl- (9CI) (CA INDEX NAME)



RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L9 ANSWER 10 OF 10 CAPLUS COPYRIGHT 2005 ACS on STN
AN 1997:533606 CAPLUS
DN 127:205350
TI Preparation of (phenoxypropylthio)phenylacetates and related compounds as antiobesity, antiatherosclerotic, and antidiabetic agents.
IN Adams, Alan D.; Doebber, Thomas W.; Berger, Joel P.; Berger, Gregory D.; Jones, Anthony B.; Von Langen, Derek; Leibowitz, Mark D.; et al.
PA Merck and Co., Inc., USA; Adams, Alan D.; Doebber, Thomas W.; Berger, Joel P.; Berger, Gregory D.; Jones, Anthony B.
SO PCT Int. Appl., 192 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 7

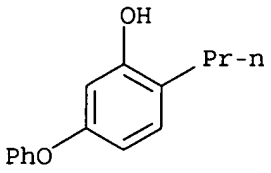
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9728115	A1	19970807	WO 1997-US1689	19970131
	W: AL, AM, AU, AZ, BA, BB, BG, BR, BY, CA, CN, CU, CZ, EE, GE, HU, IL, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LV, MD, MG, MK, MN, MX, NO, NZ, PL, RO, RU, SG, SI, SK, TJ, TM, TR, TT, UA, US, UZ, VN				
	RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
	CA 2245529	AA	19970807	CA 1997-2245529	19970131
	AU 9721159	A1	19970822	AU 1997-21159	19970131
	AU 721452	B2	20000706		
	EP 888278	A1	19990107	EP 1997-906471	19970131
	EP 888278	B1	20030723		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, FI				
	JP 2002503202	T2	20020129	JP 1997-527883	19970131
	AT 245622	E	20030815	AT 1997-906471	19970131
	ES 2202582	T3	20040401	ES 1997-906471	19970131
PRAI	US 1996-11093P	P	19960202		
	GB 1996-4231	A	19960228		
	US 1996-34435P	P	19961223		
	WO 1997-US1689	W	19970131		

OS MARPAT 127:205350
AB Title compds. [I; R = H, (substituted) alkyl, aryl, heteroaryl; R1 = H, (substituted) alkyl, alkenyl, alkynyl, cycloalkyl; R2 = H, OH, (substituted) alkyl, acyl, alkenyl, alkynyl, heteroaryl; R4 = R2, BR5, etc.; R5 = (substituted) aryl, heteroaryl; B = O, NR1, S, SO, SO2; Z = (modified) CO2H, tetrazolyl; ZW = ZCR6R7, ZCH:CH, ZCR6R7R8; R6, R7 = H, alkyl; R8 = CR6R7, O, NR6, S, SO, SO2; X1, X2 = H, OH, halo, (substituted) alkyl, alkenyl, alkynyl, aryl, aralkyl, heteroaryl, acyl, etc.; Y = S, SO, SO2, CH2, CO, CONH, O, SO2NH; Y1 = O, NR, C; Q = (unsatd.) C2-4 hydrocarbon chain], were prepared Thus, Me 3-chloro-4-dimethylcarbamoylthiophenylacetate was refluxed 2 h with NaOMe in MeOH;

the cooled solution was treated with 1-bromo-3-(2-propyl-3-hydroxy-4-propionylphenoxy)propane (preparation given) and the solution was stirred 1 h to give Me 3-chloro-4-[3-(2-propyl-3-hydroxy-4-propionylphenoxy)propylthio]phenylacetate.

IT **194793-00-5P**
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of (phenoxypropylthio)phenylacetates and related compds. as antiobesity, antiatherosclerotic, and antidiabetic agents)

RN 194793-00-5 CAPLUS
CN Phenol, 5-phenoxy-2-propyl- (9CI) (CA INDEX NAME)



=> file stnguide

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	50.30	385.38

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-7.30	-7.30

FILE 'STNGUIDE' ENTERED AT 08:26:10 ON 22 NOV 2005

FILE CONTAINS CURRENT INFORMATION.
LAST RELOADED: Nov 11, 2005 (20051111/UP).

=> SET NOTICE DISPLAY 1

NOTICE SET TO 1 U.S. DOLLAR FOR DISPLAY COMMAND
SET COMMAND COMPLETED

=> index IFICLS,PATOSEP,PATDPA,INPADOC

'PATOSEP' IS NOT A VALID FILE NAME
ENTER A FILE NAME OR (IGNORE):ignore

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.72	386.10

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-7.30

INDEX 'IFICLS, PATDPA, INPADOC' ENTERED AT 08:33:24 ON 22 NOV 2005

3 FILES IN THE FILE LIST IN STNINDEX

10/816,967

Enter SET DETAIL ON to see search term postings or to view
search error messages that display as 0* with SET DETAIL OFF.

=> log y

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.59	386.69

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-7.30

STN INTERNATIONAL LOGOFF AT 08:33:36 ON 22 NOV 2005

Connection closed by remote host